Vanguard CPV-Lepto

Authorised

- Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated
- Leptospira interrogans, serovar Canicola, strain C51, Inactivated
- Canine parvovirus, strain NL-35-D, Live

Product identification

Medicine name:
Vanguard CPV-Lepto

Active substance:
Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

Leptospira interrogans, serovar Canicola, strain C51, Inactivated

Canine parvovirus, strain NL-35-D, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Leptospira interrogans, serovar Icterohaemorrhagiae, strain NADL 11403, Inactivated

915.00 relative unit(s) / 1.00 millilitre(s)

Leptospira interrogans, serovar Canicola, strain C51, Inactivated
740.00 relative unit(s) / 1.00 millilitre(s)

Canine parvovirus, strain NL-35-D, Live
7.00 log10 50% cell culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI05

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Blister met 25 glazen (Type I) injectieflacons met rubberstop en metalen felscapsule à 1 dosis vaccin.
Blister met 100 glazen (Type I) injectieflacons met rubberstop en metalen felscapsule à 1 dosis vaccin.
Blister met 10 glazen (Type I) injectieflacons met rubberstop en metalen felscapsule à 1 dosis vaccin.
Blister met 1 glazen (Type I) injectieflacon met rubberstop en metalen felscapsule à 1 dosis vaccin.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis B.V.

Marketing authorisation date:
1/11/1994

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 8371

Date of authorisation status change:
13/12/2016

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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